



P. FORM TO-1449

U.S. DEPARTMENT OF COMMERCE
PATENT AND TRADEMARK OFFICE

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10/057,629**INFORMATION DISCLOSURE STATEMENT
BY APPLICANT***(Use several sheets if necessary)*

APPLICANT:

Harry R. Davis

FILING DATE:

January 25, 2002

GROUP:

1614**OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)**

MS	AA	Exhibit A: SCH 58235 Micronized (ezetimibe), Drug Formulation Development Summary
	AB	Exhibit B: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AC	Exhibit C: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AD	Exhibit D: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AE	Exhibit E: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AF	Exhibit F: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AG	Exhibit G: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AH	Exhibit H: SCH 58235 (ezetimibe), Drug Formulation Development Summary
	AI	Exhibit 1: Master Sheet for the SCH 58235 and Lovastatin Research Study, <i>Schering-Plough Research Institute</i> (Protocol No. C906-411), page 1576-1585
	AJ	Exhibit 2: Medical Research Study #1055/97, SCH 58235: Bioavailability of Single Oral Doses of Two Prototype Tablet Formulations and the Reference Capsule Formulation of SCH 58235 in Normal Male Volunteers: A Four Way Crossover Study #C97-221-01, Informed Consent, <i>Peninsular Testing Corporation</i> , page 106-112
	AK	Exhibit 3: Consent Form to Participate in a Research Study, "A Phase II Double Blind Dose Response Investigation of Efficacy and Safety of Four Doses of SCH 58235 Compared to Placebo in Subjects with Primary Hypercholesterolemia," <i>Schering-Plough Research Institute</i> (Protocol No. C98-010), page 1558-1566
	AL	Exhibit 4: Medical Research Study #1096/99, SCH 58235: Pharmacokinetic Pharmacodynamic Drug Interaction Study with Digoxin in Healthy Volunteers #C98-114, Informed Consent, <i>Peninsular Testing Corporation</i> , page 124-130
MS	AM	Exhibit 5: Informed Consent, "SCH 58235: Assessment of Multiple-Dose Drug Interaction Between 58235 and Gemfibrozil in Healthy Volunteers," <i>Schering-Plough Research Institute</i> , page 1-8

EXAMINER

DATE CONSIDERED

10/16/03

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.